



## AMENDMENTS TO THE CLAIMS

*Please cancel claims 16-20. Please amend the claims as follows:*

1. (Currently Amended) A tumor necrosis-factor inducible promoter, consisting of an isolated nucleic acid sequence selected from the group consisting of ~~SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6 and, SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:13, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:22, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:27, SEQ ID NO:28, and SEQ ID NO:29, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36.~~
2. (Original) An expression vector comprising one or more of the tumor necrosis-factor inducible promoters of claim 1.
3. (Original) The expression vector of claim 2 further comprising a polylinker adjacent to the 3' end of the one or more tumor necrosis-factor inducible promoters.
4. (Original) The expression vector of claim 2 comprising two or more of the tumor necrosis-factor inducible promoters.
5. (Original) The expression vector of claim 4 further comprising a polylinker adjacent to the 3' end of each of the two or more tumor necrosis-factor inducible promoters.
6. (Original) The expression vector of claim 2 further comprising a reporter gene operatively linked to the one or more of the tumor necrosis-factor inducible promoters.
7. (Original) The expression vector of claim 4 further comprising a reporter gene operatively linked to the two or more of the tumor necrosis-factor inducible promoters.
8. (Original) A recombinant host cell transfected with one or more of the expression vector of

claim 2.

9. (Original) A recombinant host cell transfected with one or more of the expression vector of claim 3.

10. (Original) A recombinant host cell transfected with one or more of the expression vector of claim 4.

11. (Original) A recombinant host cell transfected with one or more of the expression vector of claim 5.

12. (Original) A recombinant host cell transfected with one or more of the expression vector of claim 6.

13. (Original) A recombinant host cell transfected with one or more of the expression vector of claim 7.

14. (Withdrawn) A method for identifying candidate compounds for treating or preventing autoimmune disorders or cancer, comprising

- a) providing a recombinant host cell according to claim 12;
- b) contacting the recombinant host cell with tumor necrosis factor alone or together with one or more test compound under conditions suitable for expression of the reporter gene;
- c) determining reporter gene expression levels; and
- d) identifying those test compounds that modify TNF-induced reporter gene expression, wherein such modification identifies a test compound as a candidate for the treatment or prevention of autoimmunity or cancer.

15. (Withdrawn) A method for identifying candidate compounds for treating or preventing autoimmune disorders or cancer, comprising

- a) providing a recombinant host cell according to claim 13;

- b) contacting the recombinant host cell with tumor necrosis factor alone or together with one or more test compound under conditions suitable for expression of the reporter gene;
- c) determining reporter gene expression levels; and
- d) identifying those test compounds that modify TNF-induced reporter gene expression, wherein such modification identifies a test compound as a candidate for the treatment or prevention of autoimmunity or cancer.

16-20. (Cancelled)

21. (Withdrawn) A method for identifying candidate tumor necrosis factor inducible promoters, comprising:

- a) aligning a test sequence consisting of a nucleic acid sequence with a comparison sequence selected from the group consisting of ~~SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, and SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:13, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:22, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:27, SEQ ID NO:28, and SEQ ID NO:29, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36~~, using a gap opening penalty of 50 and a gap extension penalty of 3 to define a test alignment;
- b) shuffling the nucleic acid sequence of the test sequence at least one hundred times, while maintaining its length and composition, to produce a series of randomized sequences;
- c) aligning the randomized sequences with the comparison sequence using a gap opening penalty of 50 and a gap extension penalty of 3, to produce a series of randomized alignments;
- d) determining an average alignment quality of the randomized alignments, wherein the average alignment quality of the randomized alignments represents an alignment expected by chance;
- e) comparing the test alignment with the average alignment quality of the randomized alignments; and
- f) identifying a test alignment with a probability value of less than 0.05 that the alignment is obtained by chance as a candidate tumor necrosis factor inducible promoter.

22. (Withdrawn) A method for identifying candidate compounds for treating or preventing autoimmune disorders or cancer, comprising

- a) providing a recombinant host cell according to claim 12;
- b) contacting the recombinant host cell with one or more test compound under conditions suitable for expression of the reporter gene;
- c) determining reporter gene expression levels; and
- d) identifying those test compounds that modify reporter gene expression, wherein such modification identifies a test compound as a candidate for the treatment or prevention of autoimmunity or cancer.

23. (Withdrawn) A method for identifying candidate compounds for treating or preventing autoimmune disorders or cancer, comprising

- a) providing a recombinant host cell according to claim 13;
- b) contacting the recombinant host cell with one or more test compound under conditions suitable for expression of the reporter gene;
- c) determining reporter gene expression levels; and
- d) identifying those test compounds that modify reporter gene expression, wherein such modification identifies a test compound as a candidate for the treatment or prevention of autoimmunity or cancer.